patent application as follows:

Amendments to the Claims

1-16. (Cancelled)

17. (Currently amended) A compound represented by the structural formula:

$$R_4$$
 R_1
 R_2
 R_3

wherein

R₁ is hydrogen;

C₁₋₁₀ alkyl or substituted alkyl;

 $; O(CH_2)_n - Y$

 $N(COZ)(CH_2)_mY$; or

 $N[(CH_2)_mQ][(CH_2)_nY];$

 R_2 and R_3 are independently selected from:

hydrogen;

 C_{1-10} alkyl or substituted alkyl; or

 R_2 and R_3 together are cycloalkyl;

R₄ is hydrogen;

 C_{1-10} alkyl or substituted alkyl;

phenyl or substituted phenyl;

 $(CH_2)_nY$; or

 $(CH_2)_mO(CH_2)_nY;$

wherein:

m and n are independently between 1 and 10; Q and Y are independently selected from hydrogen, CO_2H or salts thereof or OPO_3^{2-} ;

Z is hydrogen or $C_{1\text{--}10}$ alkyl or substituted alkyl; and,

X is represents an amino acid, a peptide, oligopeptide or polypetitide.

18. (Currently amended) A compound represented by the structural formula:

$$R_4$$
 R_3 R_3

wherein

R₂ and R₃ are independently selected from hydrogen, C₁₋₁₀ alkyl or substituted alkyl, or R₂ and R₃ together are cycloaklyl;
R₄' is a blocking group; and,
X is represents an amino acid, a peptide;
oligopeptide or polypeptide.

19. (Previously amended) The compound of claim 18, wherein R_4 ' is selected from:

hydrogen;

 C_{1-10} alkyl or substituted alkyl; phenyl or substituted phenyl; $(CH_2)_nCO_2Y; \text{ and,}$

 $(CH_2)_n - O - (CH_2)_m Y;$

wherein:

m and n are independently between 0 and 10; and, $\mbox{$Y$ is hydrogen, or $C_{1\text{--}10}$ alkyl or substituted}$ alkyl.

20. (Currently amended) The compound of claim 17, or a salt thereof, wherein the compound is:

Methyl 1-glutaryl-7-nitroindoline-5-acetate 8,

Methyl 1-[(5-dihydroxyphosphoryloxy)pentanoyl)]
7-nitroindoline-5-acetate 9,

Methyl 1-[S-(4-amino-4-carboxybutanoyl)]-7-nitroindoline-5-acetate 10;

Methyl 1-(4-aminobutanoyl)-7-nitroindoline-5-acetate 21;

Methyl 1-acetyl-7-nitroindoline-5-acetate 16;
Mono[1-(5-methoxycarbonylmethyl-7-nitroindolyl)]

amide of 1,2-bis(O-aminophenoxy)ethane-

*N,N,N',N'-*tetraacetic acid;

1-Acetyl-4-methoxy-7-nitroindoline 25;

1-Acetyl-4-methoxy-5-methy-7-nitroindoline 30;

1-[S-(4-Amino-4-carboxybutanoy1)]-4-methoxy-7-nitroindoline;

1-(4-Aminobutanoy1)-4-methoxy-7-nitroindoline; 1-(5-Dihydroxyphosphoryloxy)pentanoy1)[-4-methoxy-7-nitroindoline;

1-[S-(4-Amino-4-carboxybutanoy1)]-4-methoxy-5-methyl-7-nitroindoline; or

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1-(4-Aminobutanoyl)-4-methoxy-5-methyl-7-
         nitroindoline. 7 or
         1-[(5-Dihydroxyphosphoryloxy)pentanoyl)]-4-
         methoxy-5-methyl-7-nitroindoline.
21.
     (Currently amended) The compound of claim 18, or a
     salt thereof, wherein the compound is:
         Methyl 1-glutaryl-7-nitroindoline-5-acetate 8;
         Methyl 1-[(5-dihydroxyphosphoryloxy)pentanoyl)]-
         7-nitroindoline-5-acetate 9;
         Methyl 1-[S-(4-amino-4-carboxybutanoy1)]-7-
         nitroindoline-5-acetate 10;
         Methyl 1-(4-aminobutanoyl)-7-nitroindoline-5-
         acetate 21;
         Methyl 1-acetyl-7-nitroindoline-5-acetate 16;
         Mono[1-(5-methoxycarbonylmethyl-7-nitroindolyl)]
         amide of 1,2-bis(O-aminophenoxy)ethane-
         N, N, N', N'-tetraacetic acid,
         1-Acetyl-4-methoxy-7-nitroindoline 25;
         1-Acetyl-4-methoxy-5-methy-7-nitroindoline 30;
         1-[S-(4-Amino-4-carboxybutanoyl)]-4-methoxy-7-
         nitroindoline;
         1-(4-Aminobutanoyl)-4-methoxy-7-nitroindoline;
         1-[(5-Dihydroxyphosphoryloxy)pentanoyl)[-4-
         methoxy-7-nitroindoline;
         1-[S-(4-Amino-4-carboxybutanoyl)]-4-methoxy-5-
         methyl-7-nitroindoline; or
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1-(4-Aminobutanoyl)-4-methoxy-5-methyl-7-

nitroindoline.; or

1-[(5-Dihydroxyphosphoryloxy)pentanoyl)]-4methoxy-5-methyl-7-nitroindoline.

- 22. (Previously added) The compound of claim 17, wherein X represents a neuroactive amino acid selected from the group of L-glutamate, GABA or glycine.
- 23. (Previously added) The compound of claim 18, wherein X represents a neuroactive amino acid selected from the group of L-glutamate, GABA or glycine.
- 24. (New) The compound of claim 17, wherein X represents

 a peptide selected from the group of thyrotrophin

 releasing hormone, an enkephalin, bradykinin or

 angiotensin II.
- 25. (New) The compound of claim 18, wherein X represents

 a peptide selected from the group of thyrotrophin

 releasing hormone, an enkephalin, bradykinin or

 angiotensin II.
- 26. (Currently amended) A composition comprising a compound of claim 17 and a pharmaceutically acceptable excipient or carrier.
- 27. (Currently amended) A composition comprising a compound of claim 18 and a pharmaceutically acceptable excipient or carrier.
- 28. (Currently amended) A composition comprising a compound of claim 19 and a pharmaceutically acceptable excipient or carrier.

- 29. (Currently amended) A composition comprising a compound of claim 20 and a pharmaceutically acceptable excipient or carrier.
- 30. (Currently amended) A composition comprising a compound of claim 21 and a pharmaceutically acceptable excipient or carrier.
- 31. (Withdrawn) A process for releasing an amino acid, a peptide or polypeptide, the process comprising irradiating a photoreleasable compound of claim 17 to cause the release of the amino acid, neuroactive amino acid, peptide, oligopeptide or polypeptide.
- 32. (Withdrawn) The process of claim 31, wherein said amino acid comprises a neuroactive amino acid.
- 33. (Withdrawn) A process for releasing an amino acid, a neuroactive amino acid, a peptide or polypeptide, the process comprising irradiating a photoreleasable compound of claim 18 to cause the release of the amino acid, peptide, oligopeptide or polypeptide.
- 34. (Withdrawn) The process of claim 33, wherein said amino acid comprises a neuroactive amino acid.
- 35. (Withdrawn) A process for releasing an amino acid, a neuroactive amino acid, a peptide or polypeptide, the process comprising irradiating a photoreleasable compound of claim 19 to cause the release of the amino acid, neuroactive amino acid, peptide, oligopeptide or polypeptide.

- 36. (Withdrawn) The process of claim 35, wherein said amino acid comprises a neuroactive amino acid.
- 37. (Withdrawn) A process for releasing an amino acid, a neuroactive amino acid, a peptide or polypeptide, the process comprising irradiating a photoreleasable compound of claim 20 to cause the release of the amino acid, neuroactive amino acid, peptide, oligopeptide or polypeptide.
- 38. (Withdrawn) The process of claim 37, wherein said amino acid comprises a neuroactive amino acid.
- 39. (Withdrawn) A process for releasing an amino acid, a neuroactive amino acid, a peptide or polypeptide, the process comprising irradiating a photoreleasable compound of claim 21 to cause the release of the amino acid, neuroactive amino acid, peptide, oligopeptide or polypeptide.
- 40. (Withdrawn) The process of claim 39, wherein said amino acid comprises a neuroactive amino acid.
- 41. (Withdrawn) A process of producing a compound of claim 17, the process comprising:
 - (a) reacting indoline or a derivatized indoline to substitute a blocking group at the 5-position;
 - (b) reacting the indoline compound of step (a) to couple an effector moiety at the heterocyclic nitrogen, the effector group having a protecting group; and,

- (c) nitrating the indoline compound of step (b) at the 7-position to produce said compound.
- 42. (Withdrawn) A process for purifying a compound of claim 17, the process comprising:
 - (a) eluting the compound from a HPLC column using aqueous methanol containing buffer salts;
 - (b) desalting fractions containing the compound obtained from step (a) on Amberlite XAD-2™ resin; and,
 - (c) eluting the resin with methanol to recover the compound.
- 43. (New) The compound of claim 17, wherein said amino acid is a neuroactive amino acid.
- 44. (New) The compound of claim 18, wherein said amino acid is a neuroactive amino acid.
- 45. (New) The compound of claim 19, wherein said amino acid is a neuroactive amino acid.
- 46. (New) A composition comprising a compound of claim 43 and a pharmaceutically acceptable excipient or carrier.
- 47. (New) A composition comprising a compound of claim 44 and a pharmaceutically acceptable excipient or carrier.
- 48. (New) A composition comprising a compound of claim 45 and a pharmaceutically acceptable excipient or carrier.